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SUMMARY OF SAFETY AND EFFECTIVENESS

In accordance with the requirements of the Safe Medical Device Act of 1990 and 21 CFR 807.92, Linvatec Corporation is hereby submitting the 510(k) Summary of Safety and Effectiveness for the 300W Xenon Light Source, 510(k) Number _____.

A. Submitter

Linvatec Corporation
11311 Concept Boulevard
Largo, Florida 33773-4908

B. Company Contact

Laura D. Krejci, RAC
Manager, Regulatory Affairs
(727) 399-5234 Telephone
(727) 399-5264 FAX

C. Device Name

Trade Name:	Linvatec 300W Xenon Light Source
Common Name:	Light Source
Classification Name:	Endoscope and accessories
Classification Number:	876.1500
Proposed Class:	Class II
Product Code:	GCT

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D. Predicate/Legally Marketed Devices

Karl Storz Endoscopy
Xenon 300W Light Source
510(k) #K962595

World of Medicine Lemke GMBH
Endoscopic Light Source XL300/L5
510(k) #K021717

E. Device Description

The Linvatec 300W Xenon Light Source is a light generating device that when used in conjunction with an endoscope will illuminate the surgical site during minimally invasive surgical procedures.

The Linvatec 300W Xenon Light Source is capable of interfacing with the ConMed I.S. Operating Room Control System to allow an alternate means for user control.

F. Intended Use

The Linvatec 300W Xenon Light Source is intended to be used with an endoscope to provide illumination during endoscopic procedures.

G. Substantial Equivalence

The Linvatec 300W Xenon Light Source described in this notification is similar in design, technology and intended use to the Karl Storz 300 Xenon Light Source (K962595) and the World of Medicine Lemke GMBH Model XL300/L5 Light Source (K021717).

The differences between the Linvatec 300W Xenon Light Source and the predicate devices are minor and raise no new questions of safety and effectiveness. Accordingly, Linvatec Corp. believes that the Linvatec 300W Xenon Light Source is equivalent to the predicate devices currently on the market.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

AUG 21 2003

Ms. Laura Krejci
Manager, Regulatory Affairs
Linvatec Corporation
11311 Concept Boulevard
Largo, Florida 33773-4908

Re: K031994
Trade/Device Name: Linvatec 300W Xenon Light Source
Regulation Number: 21 CFR 876.1500
Regulation Name: Endoscope and accessories
Regulatory Class: II
Product Code: GCT
Dated: June 26, 2003
Received: June 27, 2003

Dear Ms. Krejci:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (301) 594-4659. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address http://www.fda.gov/cdrh/dsma/dsmamain.html

Sincerely yours,

for Mark N Melkerson

Celia M. Witten, Ph.D., M.D.
Director
Division of General, Restorative
and Neurological Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure

June 26, 2003

510(k) Number (if known): K031994

Device Name: Linvatec 300W Xenon Light Source

Indications for Use:

The Linvatec 300W Xenon Light Source is intended to be used with an endoscope to provide illumination during endoscopic procedures.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE If NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use ✓ OR Over-the-Counter Use ___
(Per 21 CFR 801.109)

(Optional Format 1-2-96)

for Mark N Milhauser
(Division Sign-Off)
Division of General, Restorative
and Neurological Devices

510(k) Number K031994